# INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Not for submission under 37 CFR 1.99)

Application Number		09848774	
Filing Date		2001-05-03	
First Named Inventor	GONDA, IGOR		
Art Unit		3761	
Examiner Name			
Attorney Docket Number		AERY-058CON3	

#### CERTIFICATION STATEMENT

Please see 37 CFR 1.97 and 1.98 to make the appropriate selection(s):

That each item of information contained in the information disclosure statement was first cited in any communication [X] from a foreign patient office in a counterpart foreign application not more than three months prior to the filling of the information disclosure statement. See 37 CFR 1.97(e/11).

# OR

That no item of information contained in the information disclosure statement was cited in a communication from a foreign patent office in a counterpart foreign application, and, to the knowledge of the person signing the certification after making reasonable inquiry, no item of information contained in the information disclosure statement was known to any individual designated in 37 CFR 156(c) more than three months prior to the filing of the information disclosure statement. See 37 CFR 197(e)(c).

- See attached certification statement.
- Fee set forth in 37 CFR 1.17 (p) has been submitted herewith.

Karl Rozicevic

□ None

Name/Print

## SIGNATURE

A signature of the applicant or representative is required in accordance with CFR 1.33, 10.18. Please see CFR 1.4(d) for the form of the signature.

form of the signature.					
Signature	/Karl Bozicevic/	Date (YYYY-MM-DD)	2010-02-11		

Registration Number

28 807

This collection of information is required by 3T CFR 1.97 and 1.98. The information is required to obtain or retain a benefit by the public which is to file railed by the USPTO to process) an application. Confidentiality is governed by \$5 U.S. C. 12 and 37 CFR 1.14. This collection is estimated to take 1 hour to complete, including gathering, preparing and submitting the completed application from to the USPTO. There will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, U.S. operatment of Comments of the Comment of t

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The Privacy Act of 1974 (P. L. 95-79) requires that you be given certain information in connection with your submission of the attached form related to a patient application or patient. Accordingly, pursuant to the requirements of the Act, please be advised that: (1) the general authority for the collection of this information is 35 U.S.C. (2)(2)(2) furnishing of the information solicited to isolutionary, and (3) the principal purpose for which the information is used by the U.S. Patient and Trademan KORICs is to information, the U.S. Patient and Trademan KORICs may not be able to process and/or examine your submission, which may result in formation of proceedings or abandonment of the application or experigation of the patient.

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- A record in this system of records may be disclosed, as a routine use, to another federal agency for purposes of National Security review (35 U.S.C. 181) and for review pursuant to the Atomic Energy Act (42 U.S.C. 218(c)).
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- A record from this system of records may be disclosed, as a routine use, to the public after either publication of
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